



SmartPA Criteria Proposal

Drug/Drug Class:	NSAIDs PDL Edit		
First Implementation Date:	June 25, 2012		
Proposed Date:	December 15, 2022		
Prepared For:	MO HealthNet		
Prepared By:	MO HealthNet/Conduent		
Criteria Status:	⊠Existing Criteria □Revision of Existing Criteria □New Criteria		

Executive Summary

Purpose: The MO HealthNet Pharmacy Program will implement a state-specific preferred drug list.

Why Issue Selected:

Nonsteroidal anti-inflammatory drugs (NSAIDs) are commonly used to treat rheumatoid arthritis (RA), osteoarthritis (OA), and pain from various etiologies. NSAIDs are the most widely used drugs in the United States, with approximately 80 million prescriptions being filled yearly. These drugs, however, are associated with adverse events including gastrointestinal bleeding, peptic ulcer disease, hypertension, edema, renal disease, and increased risk of myocardial infarction.

Total program savings for the PDL classes will be regularly reviewed.

Program-Specific Information:

cific	Preferred Agents	Non-Preferred Agents
ion:	Celecoxib	Arthrotec®
	 Diclofenac 1% Gel OTC 	Celebrex®
	 Diclofenac Sodium DR/EC Tabs 	Daypro®
	 Ibuprofen 	Diclofenac 1% Gel Rx
	 Ketorolac Inj/Tabs 	Diclofenac 1.3% Patch (gen Flector®)
	 Meloxicam Tabs 	Diclofenac Topical Soln
	Naproxen OTC	Diclofenac Caps (gen Zorvolex®)
	 Naproxen Tabs Rx (gen Naprosyn®) 	Diclofenac Potassium
		Diclofenac Sodium ER (gen Voltaren®
		XR)
		Diclofenac/Misoprostol Diffusion!
		Diflunisal Duavia®
		Duexis® Floorib
		Elyxyb Etodolac
		Etodolac Etodolac ER
		Feldene®
		Fenoprofen
		Flector® Patch
		Flurbiprofen
		Ibuprofen/Famotidine
		Indocin®

	Indomethacin
	Indomethacin ER
	Ketoprofen
	Ketoprofen ER
	Ketorolac Nasal Spray
	Licart [™]
	 Lofena[™]
	Meclofenamate
	Mefenamic Acid
	Meloxicam Caps
	Mobic®
	Nabumetone
	Nalfon®
	Naprelan®
	Naprosyn®
	Naproxen CR (gen Naprelan®)
	Naproxen DS (gen Anaprox® DS)
	Naproxen EC (gen Naprosyn® EC)
	 Naproxen Sodium (gen Anaprox[®])
	Naproxen Susp
	Naproxen/Esomeprazole
	Oxaprozin
	Pennsaid®
	Piroxicam
	Qmiiz [™] ODT
	Relafen® DS
	Sulindac
	Tolmetin
	Vimovo®
	Voltaren®
☐ Increased risk of ADF	□ Preferred Drug List

Type of Criteria: ☐ Increased risk of ADE

☑ Preferred Drug List

☐ Clinical Edit

□ Databases + Prescriber-Supplied

Setting & Population

Drug class for review: NSAIDs

Age range: All appropriate MO HealthNet participants

Approval Criteria

- For non-preferred agents:
 - o For diclofenac epolamine 1.3% patch:
 - Documented diagnosis of acute pain due to minor strains, sprains or contusions in the last 30 days AND
 - Failure to achieve desired therapeutic outcomes with a trial on 2 or more preferred oral agents
 - Documented trial period for preferred agents OR
 - Documented ADE/ADR to preferred agents (gastrointestinal effects, high risk for congestive heart failure, renal failure, concomitant use of lithium) OR

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- o Failure to achieve desired therapeutic outcomes with a trial on 4 or more preferred agents
 - Documented trial period for preferred agents OR
 - Documented ADE/ADR to preferred agents AND
- For Cambia: documented diagnosis of acute migraine
- o For diclofenac sodium solution: documented diagnosis of osteoarthritis of knee
- For Elyxyb: Clinical Consultant Review for medical necessity
- o For Vimovo: documented compliance on naproxen and omeprazole single agents (30/180 days)

Denial Criteria

- Lack of adequate trial on required preferred agents
- Therapy will be denied if all approval criteria are not met
- Claim exceeds maximum dosing limitation for the following:

Drug Description	Generic Equivalent	Max Dosing Limitation
Flector 1.3% Patch	Diclofenac Epolamine	2 patches per day
Relafen 500 mg	Nabumetone	4 tablets per day
Relafen 750 mg	Nabumetone	2 tablets per day
Relafen DS 1,000 mg	Nabumetone	2 tablets per day
Sprix Nasal Spray	Ketorolac	1 bottle per day AND 5 bottles per month
Toradol 10mg tablet	Ketorolac	5 tablets per day AND 25 tablets per month
Voltaren 1% Gel	Diclofenac Sodium	17 grams per day

Required Documentation
Laboratory Results: MedWatch Form: Progress Notes: X X
Disposition of Edit
Denial: Exception Code "0160" (Preferred Drug List) Rule Type: PDL
Default Approval Period

3 months

References

- Evidence-Based Medicine and Fiscal Analysis: "NSAIDs Therapeutic Class Review", Conduent Business Services, L.L.C., Richmond, VA; November 2021.
- Evidence-Based Medicine Analysis: "Non-steroidal anti-inflammatory drugs (NSAIDs)", UMKC-DIC; August 2022.
- USPDI, Micromedex; 2022.
- Drug Facts and Comparisons On-line; 2022.